

AMENDMENTS TO THE CLAIMS

This listing of claims replaces all prior versions, and listings, of claims in the application.

1. (Currently Amended) A pharmaceutical composition, ~~comprising a non-steroidal anti-inflammatory and an opiate analgesic~~ comprising, in a unit dosage form, ketorolac and tramadol in combination with colloidal silicon dioxide, sodium glycolate starch, lactose, microcrystalline cellulose, and magnesium stearate, wherein the ketorolac is present in the composition in a range of 0.0010 g to 0.10000 g and the tramadol is present in a range of 0.0010 g to 0.20000 g, wherein the composition is in capsule form.
2. (Currently Amended) A pharmaceutical composition according to claim 1, wherein the non-steroidal anti-inflammatory is ketorolac tromethamine and the opiate analgesic is tramadol hydrochloride.
3. (Canceled)
4. (Currently Amended) A pharmaceutical composition according to ~~claim~~ claim 1, ~~wherein the ketorolac tromethamine is present in the composition in a range of 0.0010 g to 0.10000 g and the tramadol hydrochloride is present at a range of 0.0010 g to 0.2000 g, the colloidal silicon dioxide is present in a range of 0.0001 g to 0.02000 g, the sodium glycolate starch is present in a range of 0.0010 g to 0.20000 g, the lactose is present in the range of 0.0100 g to 0.50000 g, the microcrystalline cellulose is present in a range of 0.0100 g to 0.50000 g, the magnesium stearate is present in the range of 0.0001 g to 0.02000 g, and the excipient is present in the range of 0.0001 g to 1.00000 g.~~
5. (Canceled)
6. (Currently Amended) A method for preparing a composition according to claim 1 ~~2~~, comprising the following steps:
 - a) mixing ketorolac tromethamine, colloidal silicon dioxide, tramadol hydrochloride, sodium glycolate starch, lactose, microcrystalline cellulose, and magnesium stearate to produce a powdered mixture;

- b) analyzing the powdered mixture; and
- c) encapsulating and conditioning the mixture.

7. (Previously presented) A method of treating pain, the method comprising administering an effective amount of the composition according to any one of claims 1, 2 and 4-6.